

Please substitute the following set of claims for the pending claim set.

IN THE CLAIMS

1-59. (canceled)

60. (Currently amended) A hypermutable, transgenic mouse wherein the germ and somatic cells of said mouse express comprise a transgenic polynucleotide encoding a dominant negative form of a PMS2 mismatch repair protein dominant negative allele of a PMS2 mismatch repair gene, wherein said dominant negative allele comprises a PMS2 134 allele.

61. (Currently amended) A hypermutable, transgenic mouse produced by a process comprising the steps of:

introducing a transgenic polynucleotide comprising encoding a dominant negative form of a PMS2 mismatch repair protein a sequence encoding a dominant negative allele of a PMS2 mismatch repair gene into a fertilized mouse egg, wherein the dominant negative allele comprises a PMS2 134 allele, whereby said protein is expressed and said fertilized mouse egg becomes hypermutable;

implanting the fertilized egg into a pseudopregnant female; and

allowing said mouse egg to develop into a hypermutable, transgenic mouse.

62. (Currently amended) A method of making a hypermutable fertilized mouse egg comprising:

introducing into said fertilized mouse egg a transgenic polynucleotide comprising encoding a dominant negative form of a PMS2 mismatch repair protein a sequence encoding a dominant negative allele of a PMS2 mismatch repair gene, wherein the

~~dominant negative allele comprises a PMS2 134 allele~~, whereby said protein is expressed and said fertilized mouse egg becomes hypermutable.

63-70. (canceled)

71. (Currently amended) A method for generating a mutation in a gene of interest comprising the steps of:

introducing a transgenic polynucleotide ~~comprising~~ encoding a dominant negative allele form of a *PMS2* mismatch repair gene protein into a fertilized mouse egg, ~~wherein~~ the dominant negative allele comprises a PMS2 134 allele, whereby said protein is expressed and the fertilized mouse egg becomes hypermutable;

implanting the fertilized egg into a pseudopregnant female;

allowing said fertilized mouse egg to develop into a hypermutable, transgenic mouse; and

testing the mouse to determine whether the gene of interest harbors a mutation.

72. (previously presented) The method of claim 71 wherein the step of testing comprises analyzing a nucleotide sequence of the gene of interest.

73. (previously presented) The method of claim 71 wherein the step of testing comprises analyzing mRNA transcribed from the gene of interest.

74. (previously presented) The method of claim 71 wherein the step of testing comprises analyzing a protein encoded by the gene of interest.

75. (previously presented) The method of claim 71 wherein the step of testing comprises analyzing the phenotype of the gene of interest.

76-80. (canceled)

81. (Currently amended) The method of claim 62 wherein the mismatch repair gene protein is human PMS2 PMS2.

82. (Currently amended) The method of claim 81 wherein said dominant negative form of a PMS2 mismatch repair gene protein is encoded by a polynucleotide which comprises a truncation mutation at codon 134 ~~as shown in~~ of SEQ ID NO:1.

83. (Currently amended) The method of claim 82 wherein the truncation mutation is a thymidine at nucleotide 424 of wild-type PMS2 ~~as shown in~~ of SEQ ID NO:1.

84. (Currently amended) The hypermutable, transgenic mouse of claim 60 ~~comprising a~~ wherein the protein ~~which~~ consists of the first 133 amino acids of human PMS2.

85. (Currently amended) The hypermutable, transgenic mouse of claim 61 wherein the ~~mismatch repair gene~~ transgenic polynucleotide is human PMS2.

86. (Currently amended) The hypermutable, transgenic mouse of claim 61 wherein the transgenic polynucleotide ~~dominant negative allele~~ comprises a truncation mutation at codon 134 ~~of~~ ~~as shown in~~ SEQ ID NO:1.

87. (Currently amended) The hypermutable, transgenic mouse of claim 86 wherein the truncation mutation is a thymidine at nucleotide 424 of wild-type PMS2 ~~of~~ ~~as shown in~~ SEQ ID NO:1.

88. (Currently amended) The mouse of claim 60 wherein the mismatch repair gene protein is human PMS2 PMS2.

89. (Currently amended) The mouse of claim 88 wherein said transgenic polynucleotide mismatch repair gene comprises a truncation mutation at codon 134 of as shown in SEQ ID NO:1.

90. (Currently amended) The mouse of claim 89 wherein the truncation mutation is a thymidine at nucleotide 424 of wild-type *PMS2* of as shown in SEQ ID NO:1.

91. (Currently amended) The method of claim 71 wherein the mismatch repair gene protein is human *PMS2* *PMS2*.

92. (Currently amended) The method of claim 91 wherein said transgenic polynucleotide mismatch repair gene comprises a truncation mutation at codon 134 of as shown in SEQ ID NO:1.

93. (Currently amended) The method of claim 92 wherein the truncation mutation is a thymidine at nucleotide 424 of wild-type *PMS2* of as shown in SEQ ID NO:1.

94. (Currently amended) The mouse of claim 61 comprising a wherein the protein which consists of the first 133 amino acid residues of human PMS2.

95. (Currently amended) The method of claim 62 wherein the mouse egg comprises a protein which consists of the first 133 amino acid residues of human PMS2.

96. (Currently Amended) The method of claim 71 wherein said mouse comprises a protein which consists of the first 133 amino acid residues of human PMS2.